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(54) Title: DYNAMICS BIONEMS SENSORS AND ARRAYS OF BIONEMS SENSOR IMMERSSED IN FLUIDS

(57) Abstract: A bioNEMS device comprises a piezoresistive cantilever having flexing legs of which attach the cantilever to a support and a biofunctionalized portion at the tip. A bias current applied to the legs is limited by a maximal acceptable temperature increase at the biofunctionalized tip. The length of the cantilever has a magnitude chosen to minimize background Johnson noise. A catalyzed receptor on the device binds to a ligand whose binding rate coefficient is enhanced. The catalyst lowers the receptor-ligand binding activation energy and is designed by forced evolution to preferentially bind with the ligand. A carrier signal is injected by a magnetic film disposed on the cantilever which is electromagnetically coupled to a source of the carrier signal. A plurality of NEMS fluidicly coupled transducers generate a plurality of output signals from which a collective output signal is derived, either by averaging or thresholding. The NEMS devices are disposed in microfluidic flow channels and fabricated in a membrane. A linking molecule is attached to the tip of the transducer and a fluffball attached to the linking molecule to increase damping.

**DYNAMICS BIONEMS SENSORS AND ARRAYS OF BIONEMS SENSOR  
IMMERSED IN FLUIDS**

*Related Applications*

5           The present application is related to U.S. Provisional Patent Application serial  
no. 60/379,710, filed on May 7, 2002; serial no. 60/379,660, filed on May 7, 2002; serial  
no. 60/379,645, filed on May 7, 2002; serial no. 60/379,552, filed on May 7, 2002; serial  
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serial no. 60/379,643, filed on May 7, 2002; serial no. 60/379,708, filed on May 7, 2002;  
10   and serial no. 60/379,681, filed on May 7, 2002, which are incorporated herein by  
reference and to which priority is claimed pursuant to 35 USC 119.

*Incorporation of Copending Applications*

          It is to be expressly understood that the present application incorporates by  
15   reference simultaneously filed applications serial no. (PAU.34), entitled, "An Apparatus  
And Method For Vacuum-Based Nanomechanical Energy, Force, And Mass Sensors";  
and serial no. (PAU.35) entitled "A Method And Apparatus For Providing Signal  
Analysis Of A Bionems Resonator" as if set out in their entirety. Further the present  
application incorporates by reference U.S. Patent Application serial no 10/138,538, filed  
20   on May 3, 2002 entitled, "An Apparatus and Method for Ultrasensitive  
Nanoelectrochemical Mass Detection"; and U.S. Patent Application serial no.  
09/927,779, filed on Aug. 9, 2001, entitled, "Active NEMS Arrays for Biochemical  
Analyses" as if set out in their entirety.

## Background of the Invention

### 1. *Field of the Invention*

The invention relates to the field of fluidic bioNEMS devices and methods of  
5 operating the same.

### 2. *Description of the Prior Art*

There have been a number of recent advances in NEMS and in Chemical Force  
Microscopy (CFM). NEMS approaches have resulted in a family of cantilevers of small  
10 length and thickness that can resonate at high frequency with high Q. When operated  
in ideal conditions (low T, vacuum) these NEMS devices show unprecedented  
sensitivity. On a much larger size scale (AFM, CFM), work in several groups has been  
directed at analyzing the forces exerted by interactions between single molecules,  
ranging from hydrogen bonds and antibody-antigen interactions to covalent bonds.  
15 AFM cantilevers, decorated with biomolecules and interacting with derivatized surfaces  
or with derivatized magnetic beads, demonstrate forces of order 100pN for an antigen-  
antibody interaction and ~1-10nN for a covalent bond. These watershed experiments  
show the feasibility of measuring chemical events at the stochastic limit, but also  
offered evidence of the difficulty of harvesting this potential in a small, portable and  
20 robust device.

According to the invention what is needed is some way of reducing the size of  
the cantilever to NEMS dimensions to offer the needed temporal response, small  
volume and sensitivity to single molecules needed to build a device with single cell  
capability. Of course, placing a NEMS cantilever in solution and at room temperature  
25 will call for a revision of detection strategy from those usually employed for either CFM

or NEMS. The fluid will damp the NEMS cantilever making resonance detection impossible, and the thermal energy of the solution will buffet the cantilever.

What is needed is some way to exploit these to potential difficulties as an integral part of the assay.

5           In contrast to conventional CFM what is needed according to the invention is an approach which will not attempt to measure the force of a single (or small number of) chemical bond by recording cantilever deflection.

What is needed is some type of design for a NEMS cantilever will allow detection of the presence of a chemical bond by the restriction it produces in the otherwise large  
10 thermally driven motion of the cantilever using an integral sensor.

What is further needed according to the invention is a means of using an array of BioNEMS cantilevers with systematically different chemical decorations offers both high reliability and sensitivity to concentration.

Further, according to the invention what is needed is some way of interpreting  
15 the "noise" of fluctuations as signal, and biology the opportunity of assembling and employing a useful and robust assay.

Microarray technologies have provided significant recent advances in analyzing protein receptors and their ligands, as well as in analyzing gene expression profiles. For example, microarrays of a few thousand targets have become a major technique  
20 used by the drug discovery industry. These microarrays are created by photolithography, by microstamping, or by microdotting, resulting in an array of spots (20-100 $\mu$ m) on a substrate. The array is typically read by superfusing a fluorescently labeled analyte, over the array and determining the amount of binding by scanning the array with a micro-fluorimeter. Although these approaches are becoming increasingly  
25 widespread, the large size of the reader instrumentation and the intrinsic limitations of

the fluorescence analysis employed make them completely inappropriate for applications in which both portability and robust performance are required.

Furthermore, they are single-use devices, hence they cannot easily accommodate applications that require continuous monitoring. Finally, the devices rely on significant  
5 volumes of analyte, making them ill-suited to the most powerful recent advances in drug discovery provided by combinatorial chemistry or to the most sensitive assays of gene expression.

Another goal of the proposed studies is to develop a new technology of biochips at the nanoscale (BioNEMS) that is capable of sensing the binding of single biological  
10 molecules to their receptors. A growing literature of chemical force microscopy (CFM) has shown that a modified AFM can be tailored to measure the binding force of interactions ranging from single hydrogen bonds and single receptor-ligand interactions to single covalent bonds. The range of these forces are well within the capability of AFM instrumentation to detect; however, an AFM cantilever in solution does not have  
15 the temporal response characteristics needed to permit the binding and unbinding of biological ligands and their receptors to be followed reliably. Perhaps even more significant is the substantial size of the equipment required for performing AFM/CFM, and the well-known sensitivity of AFM to air-borne and surface vibrations.

What is needed is some type of technology that has the same success as CFM  
20 in detecting the forces of single molecular interactions, but is scaled down to NEMS scale to permit it to respond rapidly enough to follow the binding and unbinding events. Given the size of the chemical forces, the most robust mode for the BioNEMS to operate will be to forsake direct measurements of the force of binding. What is needed according to the invention is some type of means for using the ongoing fluctuations in

the position of the NEMS cantilever followed using integral sensors to obviate the need for the support equipment used in AFM.

### Brief Summary of the Invention

5           According to the invention the motions of the NEMS cantilevers will be used to follow binding and unbinding events. The basic idea is that a cantilever that is not coupled at its tip by a receptor-ligand pair will fluctuate in its position more dramatically than a cantilever that is restricted by a ligand-receptor pair. Strong ligand-receptor bonds can partially arrest the cantilever motion for considerable time ( $\sim t_{on}$  for the  
10   ligand-receptor pair); even weak interactions will alter the statistics of cantilever motion.

          Moving down to the small size of NEMS devices offers several significant advantages. "NEMS" in this specification is used to mean devices with at least one dimension which is equal to or smaller than one micron. It does not exclude the possibility that the "NEMS" device may have one or more other dimensions larger than  
15   one micron. Furthermore, as can be understood there is often no sharp line of distinction between the characterization of a device at or below one micron in size and one which is above one micron. The more meaningful significance to the term, "NEMS" that the device in question shares some characteristic with similar devices scaled to submicron sizes or which is unique to submicron devices or operation. As already  
20   mentioned the small size of the NEMS devices permit them to be dramatically more responsive to the kinetics of binding and unbinding. This high frequency response is critical to following the stochastic nature of receptor ligand interaction, most receptor-ligand pairs interact dynamically, binding, remaining engaged for times ranging from microseconds to seconds (depending on the exact receptor-ligand pair), and then  
25   releasing. High frequency response ( $\sim$ MHz) is critical if an assay is to follow

biomolecular interactions. The ability to resolve the opening and closing of individual membrane channels in the patch-clamp (gigaohm seal) technology has revolutionized our understanding of the physical biochemistry underlying neuronal function; before the patch-clamp, experimenters could only attempt to decode molecular mechanism from recordings of huge populations of membrane channels. It is our belief that the analysis of biomolecules is presently in exactly the same state, limited by both the vast quantities of materials required and the smearing in time inherent in even the most sensitive assays. BioNEMS are thus poised by the invention to truly move our analysis of biomolecules to the stochastic limit.

One of the powers of this approach is that it exploits the thermal motion of the cantilever, normally a major limitation in AFM, as a driving force. Furthermore, the noise of cantilever motion becomes smaller as cantilever size is decreased. In addition, the small size of NEMS devices permits an array of detectors ( $\geq 500$  cantilevers) to be constructed in a small active volume ( $\leq 100$  pL). This latter advantage is of great significance, as it offers the promise of a technology for sensing the levels of RNA, proteins and second messengers present in single cells.

The BioNEMS approach of the invention offers a major reduction in the size and nature of the instrumentation needed for it to operate (in comparison to AFM/CFM). The sensor for the motion of the cantilever will be integral to the NEMS cantilever, which eliminates the size and density limits that would be imposed by the optical detection of cantilever motion used in AFM. This will permit the BioNEMS cantilevers to be much smaller and to be dramatically more closely packed than is practical in AFM.

As outlined in the section describing NEMS sensing below, integration of a piezoresistive transducer will offer sensitivity far greater than needed to record the NEMS cantilever motions in liquid water. As a result, with proper integration, the

sensor, the detectors needed to follow cantilever motion, the logic needed to interpret the motions, and the circuitry needed to communicate the results can be packaged into a single device. In contrast to the promise of their names, current "DNA Chip" or "Proteomics Chip" technologies require bulky and heavy readers to interpret the binding of chemical species to a sensor package several centimeters in length and width. The BioNEMS approach outlined here offers the promise of package sizes consistent with the term "chip" (~DIP dimensions) and thereby offers a variety of applications that would be impossible or impractical by other approaches.

The goal of the proposed work is to exploit the thermally driven motion of the cantilever, and its modulation by receptor-ligand interactions. Knowledge of the physics of NEMS cantilevers in solution and of stochastic biochemistry will be used to interpret the motion of the cantilever. Thus, the construction of a working BioNEMS of this class requires a close collaboration between researchers working on NEMS fabrication, biologists working on the biochemical modification of device surfaces, physicists interested in the fluid dynamics of NEMS devices, and information scientists accomplished at extracting and analyzing data from arrays.

The BioNEMS research effort outlined here will have several thrusts, ranging from fundamental to applied science, and the development on new nanoscale fluidic technology. Our goal is to develop, understand and, thereby, to refine the techniques for the construction of BioNEMS and then demonstrate their novel uses.

Examples include:

- Basic studies of the performance of NEMS in solution
- Basic studies of single molecule chemistry



- Cellular studies of hormones, growth factors and second messenger. The release of growth factors from cells is typically in too low of a concentration and too small a volume for direct analysis by traditional techniques.

- Use of the BioNEMS as sensors for the output of combinatorial chemistry syntheses as a test in drug discovery efforts.

- Use as a sensitive "gene chip" for the detection of DNA sequences, or as a sensor for biological hazards.

- Use as a monitor of the concentration of environmental toxins.

The invention is defined as a submicron bioNEMS device comprising a support and a piezoresistive cantilever coupled to the support extending therefrom with a length  $l$  and having a width  $w$  and a tip, wherein the cantilever has a restriction portion of reduced width,  $b$ , and a length  $l_1$ , and a biofunctionalized portion at or near the tip. The restriction portion is comprised of multiple legs of reduced width,  $b$ , attached to the support. Preferably two legs are provided and are separated from each other by a distance of  $w - 2b$ .

The bioNEMS device further comprises a source of bias current applied to the restriction portion of the cantilever and where the magnitude of the bias current is limited by a maximal acceptable temperature increase at the biofunctionalized tip. The maximal acceptable temperature increase at the biofunctionalized tip is approximately 1 degree K.

The invention is also an improvement in a piezoresistive bioNEMS device immersed in a fluid comprising at least one oscillating cantilever having a length  $l$  having a magnitude chosen to minimize background Johnson noise relative to signal strength generated by the piezoresistive bioNEMS device. In one embodiment the signal strength is based on thermomechanical noise levels of the piezoresistive

bioNEMS device in the fluid. The piezoresistive cantilever has a width  $w$ , and a restriction portion of reduced width,  $b$ , wherein the reduced width,  $b$ , is chosen to reduce Johnson noise relative to the signal strength generated by the piezoresistive bioNEMS device.

5           The invention is further characterized as an improvement in a biofunctionalized bioNEMS device immersed in a fluid comprising a receptor disposed on the bioNEMS device for binding to a ligand of interest and a catalyst disposed on the bioNEMS device with the receptor to enhance binding rate coefficients of the receptor with the ligand of interest. The catalyst lowers the receptor-ligand binding activation energy. In  
10 one embodiment the receptor is designed by forced evolution to preferentially bind with the ligand of interest.

          The invention is also defined as a submicron device comprising a source of a carrier signal, a support, a piezoresistive cantilever coupled to the support and extending therefrom, and an element disposed on the cantilever and  
15 electromagnetically coupled to the source so that the cantilever is driven by the carrier signal from the source. The element comprises a magnetic film disposed on the cantilever, and the source generates an electromagnetic signal coupled to the magnetic film.

          Still further the invention is an apparatus comprising a plurality of NEMS  
20 resonators or transducers, each of the NEMS transducers generating an output signal; and means or circuit for processing the plurality of corresponding output signals from the plurality of NEMS transducers to obtain a collective output signal. The means averages the plurality of output signals so that the collective output signal is an average. The means determines if a predetermined fraction of the plurality of output  
25 signals are above a threshold within a predetermined time window. Each of the

plurality of NEMS transducers is biofunctionalized and the means effectively increases ligand capture rates by only generating a collective output signal indicative of an increase ligand capture rate as compared to a single one of the NEMS transducers.

The invention is described as an apparatus operating in a fluid comprising a plurality of NEMS transducers immersed in the fluid forming an array of adjacent transducers, each of the NEMS transducers generating an output signal, the motion of two adjacent NEMS transducers being coupled to each through the fluid in which the adjacent NEMS transducers are immersed. A cross correlation of movement of a first and second NEMS transducer exists and comprises the two adjacent NEMS transducers,  $C_{12} = \langle x_1(0) x_2(t) \rangle$ , as defined by

$$\frac{d}{dt} C_{12}(t) = -k_B T X_{12}(t) \quad \text{for } t > 0$$

where  $k_B$  is the Boltzman constant,  $T$  is the temperature of the fluid,  $t$  is time, and  $X_{12}$  the "susceptibility" giving the displacement  $x_2(t)$  of the second transducer for a force  $F_1$  acting on the first transducer, such that an ensemble average for the position of the second transducer is defined by

$$\langle x_2(t) \rangle = \int_{-\infty}^{\infty} X_{12}(t-t') F_1(t') dt'.$$

The invention is an apparatus operating in a fluid comprising a microfluidic flow channel for carrying a flow of the fluid, and at least one NEMS transducer disposed in the microfluidic flow channel so that a characteristic of the fluid is sensed by the NEMS transducer. The NEMS transducer is biofunctionalized and the characteristic of the fluid is sensed by the NEMS transducer is the presence or absence within the fluid of a ligand to which the NEMS transducer has been biofunctionalized.

The apparatus further comprises a plurality of the NEMS transducers, each of which is disposed in common in the flow channel. The apparatus further comprises a

plurality of flow channels among which the plurality of NEMS transducers re distributed.  
The plurality of NEMS transducers are surface fabricated or membrane fabricated.

The invention includes a method of fabricating a bioNEMS device from a  
membrane comprising the steps of providing a heterostructure comprising a wafer  
5 layer, an etch stop layer on the wafer layer, a NEMS device layer on the etch stop  
layer, and a piezoresistive layer on the NEMS device layer. Trenches are etched  
through the wafer layer to the etch stop layer to define an area which will become a  
membrane in which the NEMS device will be defined. The etch stop layer is removed  
from the bottom of the trenches to the device layer to form the membrane. Conductive  
10 contacts are selectively formed on the piezoresistive layer of the membrane by electron  
beam lithography. Regions are selectively formed which will become biofunctionalized  
on the piezoresistive layer of the membrane by electron beam lithography. A NEMS  
device is selectively formed on the piezoresistive layer of the membrane by electron  
beam lithography which include the region which will become biofunctionalized. The  
15 membrane is selectively plasma etched to remove unmasked portions to define a  
suspended NEMS device. A flow channel is selectively molded in an elastomeric layer  
disposed around the membrane. Selected regions are biofunctionalized on the NEMS  
device.

The step of providing a heterostructure further comprises polishing the wafer  
20 layer to promote adhesion to the elastomeric layer and thinning the wafer layer. The  
step of selectively plasma etching the membrane to remove unmasked portions to  
define a suspended NEMS device comprises selectively vertically plasma etching away  
unmasked portions of the NEMS device layer. The step of selectively molding a flow  
channel in an elastomeric layer disposed around the membrane comprises selectively  
25 disposing a photoresist layer to define the flow channel, disposing an elastomeric layer

on the selectively disposed photoresist layer, and removing the photoresist layer to define the flow channel.

The invention discloses a NEMS device for operating in a fluid comprising a resonating member having a tip immersed in the fluid, a linking molecule attached to the tip; and a fluffball attached to the linking molecule and providing a damping force to  
5 dissipate noise applied to the member from the fluid.

The invention also includes methods for operating the NEMS devices disclosed above.

While the apparatus and method has or will be described for the sake of  
10 grammatical fluidity with functional explanations, it is to be expressly understood that the claims, unless expressly formulated under 35 USC 112, are not to be construed as necessarily limited in any way by the construction of "means" or "steps" limitations, but are to be accorded the full scope of the meaning and equivalents of the definition provided by the claims under the judicial doctrine of equivalents, and in the case where  
15 the claims are expressly formulated under 35 USC 112 are to be accorded full statutory equivalents under 35 USC 112. The invention can be better visualized by turning now to the following drawings wherein like elements are referenced by like numerals.

### **Brief Description of the Drawings**

20 Fig. 1 is a graph of the observed improvement in the strength of the fluid coupled thermomechanical noise relative to the Johnson noise with all other dimensions fixed.

Fig. 2 is a graph of the fluid damped thermomechanical noise.

Fig. 3 is a graph of the expected signal in several liquids for a bias current of  $250\mu A$ , for a cantilever with  $b=0.6\mu m$ ,  $t=130nm$ ,  $w=2.5\mu m$ ,  $l=15\mu m$  and  $l_1=0.6\mu m$ .

25 Fig. 4 is a diagrammatic side elevational view of the cantilever in microscopically

enlarged scale.

Fig. 5 is a graph of the fractional receptor occupancy as a function of nonspecific binding events.

Fig. 6 is a diagram of a detector known as a "phase-detector" or "lock-in amplifier", or a correlation receiver.

Fig. 7 is a graph illustrating detector performance  $P_d$  as a function of SNR.

Fig. 8a is a diagrammatic top plan view of a double cantilever system.

Fig. 8b is a diagrammatic side cross-sectional view of the system of Fig. 8a.

Fig. 8c is a diagrammatic top plan view of another embodiment wherein the cantilevers are ligand coupled.

Fig. 9 is a graph of the fluid velocity component parallel to an oscillating cantilever as a function of distance,  $r$ .

Fig. 10 is a cross sectional schematic side view of a piezoresistive cantilever coupled to a microfluidic flow channel

Figs. 11a – 11c is a diagrammatic perspective view in increasing magnification of the array of cantilevers shown in Fig. 10

Fig. 12a is a scanning electron microscopic photograph of a bioMEMS transducer. Fig. 12b is a diagram of the top plan view of the transducer of Fig. 12a.

Figs. 13a – 13m are a series of diagrams illustrating a method for fabricating a bioNEMS fluidic sensor from a membrane.

Fig. 14 is a diagram modeling the dynamic action of a cantilever with a molecularly bound fluffball as an added damper.

The invention and its various embodiments can now be better understood by turning to the following detailed description of the preferred embodiments which are presented as illustrated examples of the invention defined in the claims. It is expressly

understood that the invention as defined by the claims may be broader than the illustrated embodiments described below.

### Detailed Description of the Preferred Embodiments

5           The illustrated embodiment is directed to the constraints upon the level of current bias that can be applied to a piezoresistive BioNEMS device in fluid. The force sensitivity attainable clearly hinges on the maximum level of current bias that is tolerable, given that the responsivity is proportional to bias current,  $R = I/G$ . The largest practical level of bias current is determined by the maximum temperature rise in the  
10   BioNEMS which is deemed acceptable.

          For the purposes of illustration the bioNEMS transducer or cantilever 10 shown in perspective view in the microphotograph Fig. 12a and in the top plan view diagram of Fig. 12b shall be assumed, which can be analogized as having the form of "a diving board with a cutout at its base". However, it is to be expressly understood that the  
15   geometry of the transducer 10 is to be entirely general and includes any type cantilever, doubly clamped beam, paddle or any other submicron oscillating structure. The geometry of the device 10 causes maximization to occur predominantly within a constriction region 12 comprised of one or more legs 20 of width  $b$ , as shown in Fig. 12b which region 12 allows for enhanced or variably designed flexural stiffness of  
20   cantilever 16 without restricting the fluidic damping characteristics of cantilever 16 dependent on its over length  $l$  and width  $w$ . It is also to be understood that cantilever 16 will have conventional electrodes (not shown) provided whereby a conventional external measurement circuit (not shown) providing a bias current may measure the change in piezoresistivity of legs 20 as they flex. In addition, an external driving force  
25   may or may not be applied in a conventional manner to cantilever 16 depending on the

application and design choice.

In the preferred embodiment there are two legs 20. We assume that a temperature rise of order 1 K is tolerable at the biofunctionalized tip 14 of the cantilever 16, which has a length,  $l$ , a width,  $w$ , and a thickness,  $t$ , resonant frequency in vacuum  $\omega_0/2\pi$  and force constant  $K$ .

We treat the problem as one dimensional, with the constricted region a beam 16, of length  $l_1$  and cross sectional area  $A$ , heat sunk at the supporting end 17 to supporting substrate 18. In the illustrated embodiment, cantilever 16 is made of silicon and assumed to be immersed in water, but any nanomachinable material may be employed and any ambient fluid may be contemplated. The distance  $x$  is measured from the connection of cantilever 16 to support 17 towards its tip 14. For  $x > l_1$ , a rough estimate of the heat loss to the water or fluid in which device 10 is immersed may be obtained

through the relationship  $\kappa_{Si} A \frac{d^2 T}{dx^2} = \kappa_{H_2O} P \nabla T$  where  $P$  is the perimeter around cross-sectional area  $A$  of the beam 16. Estimating  $\nabla T \sim T/w$ , and  $\frac{d^2 T}{dx^2} \sim \frac{2(w+t)\kappa_{H_2O}}{\kappa_{Si} t w^2}$  where

$\kappa_{Si} = 1.48 \times 10^2$  W/mK is the thermal conductivity of silicon and  $\kappa_{H_2O} = 0.607$  W/mK is the thermal conductivity of water. In the dissipative region  $x < l_1$  we have

$$\kappa_{Si} t b \frac{d^2 T}{dx^2} \sim -I^2 R + 4(b+t) \frac{T}{b} \kappa_{H_2O}. \text{ As boundary conditions, we have that the temperature}$$

is continuous at  $l_1$ , as is the heat flux; and the temperature must monotonically decrease for  $x > l_1$ .

We consider the three illustrative devices 10 from Table 1. For the first cantilever 16, this simple thermal conductance calculation indicates that a 1 K temperature rise at the biofunctionalized tip 14 is attained with a steady-state bias



current  $I = 250\mu\text{A}$ , leading to a power dissipation of roughly  $10,670\mu\text{W}$ . The maximal temperature rise of  $12\text{K}$  occurs within the constricted region 18, approximately  $2.3\mu\text{m}$  from the edge of substrate 18. For this bias current, the first device yields a responsivity  $R = I/G$  of about  $8\mu\text{V/nm}$ .

- 5 For the second cantilever 16 in Table 1 we allow the same  $12\text{K}$  maximal temperature rise in the constricted region 12, this coincides with a temperature rise of  $0.04\text{K}$  at the tip 14 and occurs for a current of  $75\mu\text{A}$ . For this device 10 we expect a gauge factor  $G = 5.2 \times 10^9 \Omega/\text{m}$ . Hence the expected responsivity is  $390\mu\text{V/nm}$ .

- 10 Finally, for the third cantilever 16 in Table 1, using a  $12\text{K}$  maximal temperature rise in the constricted region 12 and  $0.04\text{K}$  at the tip 14, we can allow a current of  $22\mu\text{A}$ . For this device we anticipate  $G = 5.3 \times 10^{10} \Omega/\text{m}$ . Hence the expected responsivity is  $1.2\text{mV/nm}$ .

Table 1

15

#	$t$	$w$	$\ell$	$\ell_1$	$b$	$\omega_0 / 2\pi$	$K$
1	130nm	$2.5\mu\text{m}$	$15\mu\text{m}$	$4.0\mu\text{m}$	$0.6\mu\text{m}$	0.51MHz	34mN/m
2	130nm	300nm	$10\mu\text{m}$	$2.0\mu\text{m}$	100nm	1.3MHz	20mN/m
3	30nm	100nm	$3\mu\text{m}$	$0.6\mu\text{m}$	33nm	3.4MHz	3.0mN/m

### Scaling of Fluidic Coupling in BioNEMS

- 20 For applications in which device 10 is driven by fluid-coupled thermomechanical noise in the fluid in which it is immersed, it is beneficial that this signal be maximized relative to the background Johnson noise in device 10. The relative strength of these

two sources of thermal noise is determined by the cantilever's 16 dimensions. These dimensions enter both into the magnitude of the fluidic damping which determines the spectral density of the fluid-coupled thermomechanical noise in the force domain and into the response function of the cantilever 16 (through the damping, effective mass, gauge factor, spring constant and allowed current for the same amount of heating at the tip).

For fixed thickness,  $t$ , a substantial improvement in signal strength may be obtained by decreasing the width of the cantilever legs 20 as illustrated in the graph of Fig. 1 where measured signal strength of device 10 in  $\text{nV/Hz}^{1/2}$  is graphed against frequency of the thermomechanical noise and Johnson noise for devices with  $b = 0.4\mu\text{m}$  and  $b = 0.1\mu\text{m}$ . Increasing the total length  $l$  and width  $w$  of the cantilever 16 increases the coupling to the fluid, increasing both the fluidic damping and the cooling efficiency; also leading to an improved signal-to-noise ratio.

The effect of increasing the length  $l$  of cantilever 16 is shown in the graph of Fig. 2, for cantilevers 16 of different lengths  $l$  as shown in the tabular inset with  $b = 0.1\mu\text{m}$ ,  $t = 130\text{nm}$ , and  $w = 5\mu\text{m}$  in common among each example. The ambient fluid was diethylene glycol. Fig. 2 is a graph of the fluid damped thermomechanical noise, which is predicted to increase relative to the background Johnson noise as the cantilever length is increased. For a length of  $35\mu\text{m}$ , the fluid damped thermomechanical noise has a peak an order of magnitude larger than the background Johnson noise. For each cantilever length, the bias current was chosen such that the estimated temperature rise at the tip 14 not exceed  $1^\circ\text{C}$  and that at the point of maximal temperature in the legs 20 did not exceed  $50^\circ\text{C}$ . The bias currents used and estimated temperature rises are summarized in Table 2.

Table 2

$\ell$	I	$\Delta T_{\text{tip}}$	$\Delta T_{\text{max}}$
15 $\mu\text{m}$	115 $\mu\text{A}$	1°C	8°C
18 $\mu\text{m}$	140 $\mu\text{A}$	1°C	12°C
20 $\mu\text{m}$	160 $\mu\text{A}$	1°C	16°C
25 $\mu\text{m}$	205 $\mu\text{A}$	1°C	26°C
35 $\mu\text{m}$	285 $\mu\text{A}$	0.6°C	50°C
Diethylene Glycol			
35 $\mu\text{m}$	140 $\mu\text{A}$	1°C	15°C

5 In addition to the dependence on the cantilever dimensions, it should also be noted that the magnitude of the fluid-coupled thermomechanical noise is dependent on the fluid viscosity. Fig. 3 shows the expected signal in the voltage domain for the Johnson noise and thermomechanical noise for a cantilever 16 with  $b = 0.6\mu\text{m}$ ,  $t = 130\text{nm}$ ,  $w = 2.5\mu\text{m}$ ,  $l = 15\mu\text{m}$  and  $l_1 = 0.6\mu\text{m}$  and a bias current of 250  $\mu\text{A}$  for fluids with

10 a range of different viscosities corresponding to water, diethylene glycol, glycerol, and ethylene glycol.

#### *Enhancing Receptor-Ligand Reaction Probabilities*

In one embodiment, device 10 is biofunctionalized by having bioreceptor

15 molecules 24 attached to or near tip 14 of cantilever 16 as diagrammatically shown in Fig. 4. Since one of the fundamental features of NEMS device 10 is the use of biological receptor molecules 24 attached to a "functionalized" region of an elastic cantilever 16, it is important that the receptor-ligand binding reaction have the highest

possible rate coefficient (reaction probability) for the target ligand 22 of interest.

However, it is well known that biologically expressed receptor molecules 24 do not in general have the highest possible binding rate coefficients for their target ligands, since evolution does not select for this attribute. Thus it will be useful to examine various  
5 methods for enhancing the binding rate coefficients for receptors 24 of interest in applications of the NEMS device 10.

One possible approach is to use the equivalent of catalyzed reaction by finding specific molecules which, when in close association with a particular specie of receptor 24, lower the receptor-ligand binding activation energy. Since rate coefficients, and thus  
10 reaction probabilities, are very sensitive to activation energy, e.g. have an exponential dependence, lowering this energy by even relatively small amounts yields large increases in the binding rate coefficients. Thus, the idea here is to first attach a layer of these "catalyst" molecules 26 to a small functionalized region 28 of the cantilever 16, and then attach the receptors 24 specific to the target ligand 22 of interest. The basic  
15 concept of this approach is shown in Fig. 4. The choice a catalyst in any given instance would be dictated by the ligand-receptor pair according well known principles.

A second and probably viable approach is to design the receptor molecules 24 whose binding rate coefficient is maximized with respect to a specific ligand 22 of interest. The term, "design" is understood here to mean that one uses forced evolution,  
20 biochemical techniques to select for genes that express the desired receptor 24 with higher and higher binding affinities for the chosen ligand 22 as one goes through multiple rounds of "evolving" the gene. This is usually accomplished by inserting multiple copies of the gene for a receptor 24 of interest into the genome of a particular bacterium so that the bacteria will express this receptor on their cell surfaces.

25 Binding affinity assays are then carried out on the first generation of bacteria,

and only those with the highest binding affinities are kept for the next round of the "evolutionary" cycle. Prior to starting the next cycle, variations of the receptor genes are created either through point mutations, or more efficiently by "DNA-shuffling". These cycles are then repeated until it becomes clear that no further increase in binding affinity is possible. One might expect to achieve at least an order of magnitude increase in the binding rate coefficients for particular receptor-ligand pairs by utilizing this technology.

#### *Non-Specific Ligand Binding Effects*

Fig. 5 is a graph which shows an example of the possible effects of "background" non-specific binding on the response of NEMS cantilever 16. The curve 30 is the fractional occupancy of functionalized sites 28 by target ligand 22, while curve 32 represents the fractional occupancy by both target 22 and background ligand molecules 32. The conditions used were: (1) 1000 target ligand molecules 22; (2) 100 receptor molecules 24; and (3) 10,000 background ligand molecules 32; with a non-specific binding affinity of 200 times less than for the target ligand molecules 22. The effects of non-specific binding competition for receptors 24 could be significant for device 10 as shown in Fig. 5.

#### *Use of Multiple Cantilevers for Signal Generation in BioNEMS*

The case of single undriven cantilevers 16 for biomolecule detection has been analyzed in detail copending patent application entitled, A METHOD AND APPARATUS FOR PROVIDING SIGNAL ANALYSIS OF A BIONEMS RESONATOR, serial no. (CIT.PAU.35)\_\_\_\_\_, simultaneously filed herewith and which is incorporated herein by reference. There we analyzed the expected detection performance for

several driven two-cantilever systems. In the illustrated embodiment here we explore some general issues related to the use of multiple cantilevers in BioNEMS devices 10. We approach this analysis from the standpoint of signal "design"; which is to say that we are looking for device configurations that will produce the optimum signal with  
5 respect to detection performance. Such a signal is given by

$$r(t) = A\cos(\omega_0 t + \theta) + n(t), \quad 1.1$$

where  $A$  is the amplitude of the signal, oscillating with frequency  $\omega_0$ ; and  $n(t)$  is a  
10 zero-mean Gaussian noise process with variance  $\sigma_n^2$ . Now, if both  $A$  and  $\omega_0$  are deterministic, *i.e.*, not subject to fluctuations, then the optimum detector is known to be the so called "phase-detector" or "lock-in amplifier", a block diagram of which is shown in Fig. 6. In communications theory such a detector is called a correlation receiver. Thus, we will look at device configurations that allow us to inject a "carrier" signal, at a  
15 frequency of  $\omega_0$ , such that target ligand binding events will "modulate" this carrier, *i. e.*, modify the value of  $A$ . The correlation detector of Fig. 6 is comprised of a narrowband filter 34 which takes its input,  $r(t)$ , from device 10. A reference signal from oscillator 36 is mixed by mixer 38 with the output of filter 34. The mixed filtered signal is input into low pass filter 40 and then coupled to a thresholding and decision circuit 42, which  
20 determines first if the signal is qualified as a valid or information-bearing signal and then if qualified a decision algorithm is implemented to determine if device 10 has or has not detected the ligand-receptor interaction of interest. These circuits may be implemented in analog or digital signal processors or computers as controlled by hardware design, firmware or software as devised by conventional design options.

25 We must deal with a relatively harsh constraint however, which is the severe loss

of performance, if either the amplitude,  $A$ , or the phase,  $\theta$ , of the signal are subject to significant random fluctuations. This loss of performance is illustrated in Fig. 7, where Case A is the "no random fluctuation" case; Case B is where there are random amplitude fluctuations; and Case C is the case where there is loss of phase information through random fluctuations. The case labeled "Chopper" is the performance of a single, undriven cantilever 16 using stochastic detection with multiple-sample summation as disclosed in the above incorporated referenced copending application. The method of injecting a carrier signal and "coupling" it to ligand-receptor binding events thus becomes a material consideration to obtain meaningful measurements.

10 Figs. 8a – 8c show two possible embodiments that can generate signals of the form of Eqn. 1. In Fig. 8a carrier injection is achieved by mechanically driving an unfunctionalized cantilever 16a with no receptors at a fixed frequency,  $\omega_0$ , and amplitude. The functionalized cantilever 16b responds to the driver 16a through fluid dynamic coupling of the fluid in which the device 10 is immersed. The driven cantilever 15 16b has a piezoresistive portion 46. Both cantilevers 16a and 16b are connected to supports 48, which in turn may be connected to substrates 44 or 50.

In Figs. 8a – 8c the "no signal" case is where the cantilever 16b is constrained not to move significantly by "tacking" it to the substrate 44; the presence of "free" target ligand 22 results in breaking this condition through competitive binding. Note that the side-by-side arrangement of Fig. 8a is not the preferred configuration. As a practical matter it is better to have the two cantilevers opposed, *i.e.*, tip-to-tip. Fig. 8c is a slightly different version of Figs. 8a and 8b, where the no-signal state has the two cantilevers 16 and 34 mechanically coupled through a ligand-receptor binding arrangement. Again, presence of free target ligand disrupts this state through competitive binding, resulting 25 in loss of a coherent signal.

Although we are discussing multiple cantilevers 16a and 16b, it should be pointed out that we can achieve very efficient carrier injection using a single cantilever 16 which has a small area 28 plated with a magnetic film; the external carrier signal source 36 is then generates a driving signal and is magnetically coupled to the  
5 cantilever 16 to obtain "constant" amplitude, constant phase oscillation of the cantilever beam 16. In fact, this approach probably allows us to satisfy the parameter fluctuation constraint better than any other. The coupling may also be extended to electrostatic couplings using a dipolar or paraelectric film.

A different use of multiple cantilevers 16 is to utilize an array of N identical  
10 undriven cantilevers in either an averaging mode or a coincidence mode. In the averaging mode we simply use the N outputs to obtain a  $\sqrt{N}$  improvement in our estimate of the variance. In coincidence mode, we use the constraint that some fraction of the N outputs must be above a threshold value within a fixed time window to be considered a "signal present" event. In both cases the method of using multiple  
15 cantilevers seeks to improve the ligand capture rate by increasing the number of available receptors 24.

#### *Reducing Fluidic-Induced Correlated Fluctuations*

Although the study of the single cantilever 16 in the fluid is quite well developed,  
20 there has been little attention paid to the fluid coupling of arrays of cantilevers 16. The fluid disturbance produced by the vibrating cantilever 16 is long range, falling off only as the inverse power of the separation distance between cantilevers 16. Moving one cantilever 16 will produce motion in the other cantilevers 16 through the viscous drag, or a fluidic coupling between the two cantilevers. There will correspondingly be a  
25 correlation of the stochastic motion of the cantilevers 16, since the stochastic force due



to the molecular collisions on one cantilever 16 will induce motion of the second cantilever 16 through this coupling.

This connection is made quantitative through fluctuation dissipation theory. The fluidic-induced correlated fluctuations will tend to obscure the correlations induced by a molecule tethered between the cantilevers 16, making the detection and  
5 characterization of biomolecules using this scheme harder. It is important therefore to understand the fluidic induced correlations in the absence of biomolecules, and design geometries and protocols to minimize them.

Recent experiments by Meiners and Quake on laser-tweezer controlled spherical  
10 beads provide a first indication of the results to be expected. See J. C. Meiners et.al., *Direct Measurement Of Hydrodynamic Cross Correlations Between Two Particle In An External Potential*, Phys. Rev. Lett. 82, 2211 (1999). These two authors studied the correlations induced by the fluid interaction of two 1 $\mu$ m latex beads with separation, about 3 - 10 $\mu$ m. They find a strong anticorrelation of the spheres with a maximum  
15 anticorrelation approaching the mean square displacement of a single sphere for the closest separation (3 diameters) that they investigated. This suggests a strong fluidic coupling that is hard to eliminate. However, the flow around a long cantilever 16 has quite different properties than the flow around a sphere, and this suggests a strategy for minimizing the fluidic correlations.

20 The low Reynolds number flow around a moving sphere is everywhere in phase with the motion of the sphere and falls off as 1/r. A simple model of our cantilevers 16 is to approximate them as cylinders, long compared to the radius. In this case the Stokes results for the flow around infinite cylinders can be used. The flow around an oscillating cylinder is more complicated than for spheres There is, in fact, no solution  
25 independent of Reynolds number for small values of the Reynolds number, and there

are nontrivial frequency and distance dependent phase relationships between the velocity of the fluid and the velocity of the cylinder, varying on a distance scale of a few times  $aR^{-1/2}$ . Here  $a$  is the cylinder radius which we set to the cantilever width, and  $R = \omega a^2/4\nu$  is the Reynolds number, with  $\omega$  the frequency and  $\nu$  the kinematic viscosity of the fluid.

For BioNEMS cantilevers 16,  $R$  is typically about unity. This velocity field is responsible for the motion of the second cylinder, so that there is a separation and frequency dependence to the motion of the second cylinder induced by the force on the first cylinder. Through the fluctuation-dissipation theorem, which states the fluctuations present in a system is proportional to the damping in the system, this gives us a separation and frequency dependence to the noise correlation. A plot of the velocity field as a function of the distance  $r/a$  from the cylinder axis in Fig. 9 shows in fact that the different quadratures of the velocity field have null points at different distances, suggesting that it might be possible to find parameters (cylinder separation/radius and frequency) corresponding to null points in the one or other phase of the fluid induced correlated noise. Fig. 9 shows the fluid velocity component parallel to the velocity of an oscillating cantilever 16 as a function of the distance  $r$  in units of the cantilever width  $a$  for a Reynolds number  $R = 1$ . Note the nontrivial phase relationship given by the real (in phase) and imaginary quadrature components, and the null points of the different quadrature components.

A calculation of the correlations allowing design optimization proceeds as follows. The precise relationship between the cross correlation  $C_{12} = \langle x_1(0) x_2(t) \rangle$  between the displacements of the two cantilevers and the deterministic fluidic coupling between the cantilevers is

$$\frac{d}{dt}C_{12}(t) = -k_b T X_{12}(t) \quad \text{for } t > 0 \quad 4.1$$

where  $X_{12}$  is the "susceptibility" giving the displacement  $x_2(t)$  of the *second* cantilever 16 for a force  $F_1$  acting on the first cantilever 16

5

$$\langle x_2(t) \rangle = \int_{-\infty}^{\infty} X_{12}(t-t') F_1(t') dt' \quad 4.2$$

The angular brackets denote an ensemble average, emphasizing again that a  
 10 deterministic estimate, calculation, or experiment enables the stochastic motion to be quantified. The susceptibility  $X_{12}$  can be calculated from the Stokes velocity field plotted in Fig. 9, and so the noise correlations can be predicted, although not presented here.

#### *Merging NEMS to Microfluidics*

15 Fig. 10 shows a schematic of a cross section of a device 10 with a cantilever 16 coupled to a microfluidic flow channel 52. The region etched through the wafer 54 forms part of the final flow channel. An entire array of cantilevers 16 can be fabricated within a single flow channel 54.. It is also possible to have multiple channels 54 with different devices 10 in different channels 54 depending on the desired application. It is also  
 20 possible to surface fabricate the flow channels 54 directly in the silicon. This is shown schematically for an array 56 of devices in Figs. 11a – 11c which are three perspective views in increasingly enlarged scale showing a plurality of cantilevers 16 supported on parallel supports 48 forming rows of parallel cantilevers 16 in a single flow chamber 58 communicated with an inlet and outlet flow channel 54.

*Fabrication of a BioNems Membrane Based Device*

Figs. 13a – 13g show a flow diagram schematically depicting the steps involved in the fabrication of membrane-based BioNEMS devices 10. Fabrication of these devices begins with the side cross-sectional view of Fig. 13a with a silicon device layer 148 on insulator wafer 152, 154, such as a 375 nm SiO<sub>2</sub> layer 152 on a 675μm Si layer 154. The buried oxide layer 152 must be thick enough to serve as a stop layer in the etching step through the back of the wafer 154. The Si device layer 148 should be of the desired thickness for the undoped portion of the silicon cantilever between 20nm and 100nm for most of the devices under consideration. In the illustrated embodiment an 80nm Si layer 148 is disposed beneath a 30nm heavily doped Si layer 150. The resistivity of layer 148 should be high relative to that of the heavily doped layer 150 which will be grown. (10Ωcm is sufficient).

The backside of the wafer 154 is polished. This is necessary for the adhesion of an elastomeric material thereto in which microfluidic channels 52 will be defined as described below. The wafer 154 may be thinned at the same time which serves both to decrease the ultimate volume of the flow channel 52 and reduce the necessary thickness through which one must etch in a later etching step. A final wafer thickness of 300μm is reasonable to maintain structural integrity of the wafer 154 while reducing the unnecessary material.

Next a layer of heavily boron doped silicon layer 150 is epitaxially grown on the top surface of layer 148, which layer 150 will form the conducting layer of the piezoresistor which forms part of the NEMS device 10. For most of the devices 10 this layer 150 is between 7nm and 30nm thick. The resistivity must be low compared to that of layer 148 below. A doping level of  $4 \times 10^{19} \text{ cm}^{-3}$  is typical.

The next step involves the fabrication of membranes by etching through the back of the wafer 154 as shown in the side cross-sectional view of Fig. 13b. This may be performed using a Bosch deep reactive ion etch (DRIE) to form trenches 158 through layer 154. For this step the use of a photoresist or oxide mask 156 of approximately six  
5 microns is sufficient as shown in Fig. 13b. Membranes of  $50\mu\text{m}^2$  have been used, but this is arbitrary and the dimensions to be used are determined by the application. The oxide layer 152 directly under the silicon membrane 148 is then removed from the bottom of trenches 158 with hydrofluoric acid as shown in Fig. 13c to define an area which will become membrane 162.

10 Photolithography and metal deposition is next performed on the top side of the device 10 on layer 150, aligned to the membranes 162, to form the contact pads 160 as depicted in the top plan view of Fig. 13d which shows a plurality of dies being simultaneously formed. 30nm of chrome (as an adhesion layer) followed by 250 nm of Au are deposited in the desired pattern for the bond pads 160 to form ohmic contacts to  
15 the boron doped silicon layer 150.

Silicon nitride layer 174 (300nm) is then deposited over the Au followed by 200nm silicon dioxide layer 175 to passivate the devices 10 as seen in the top plan view of Fig. 13e followed by a layer 176 of chrome as seen in the top plan view of Fig. 13f to protect the silicon dioxide 175 during the fabrication process and provide  
20 electrical continuity across the step height formed by the passivating layers as shown in Fig. 13g.

Electron beam lithography is then used to pattern first the gold pads (not shown in drawings) at the tips 14 of where the cantilever 16 will be biofunctionalized and then the cantilevers 16 are patterned (on PMMA) followed by the evaporation of a 30nm  
25 layer 178 (not shown in drawings) of chromium and liftoff as shown in the perspective

view of Fig. 13i. This portion of the fabrication process involves two steps. First there is the step of disposing a gold square 180 (not shown in drawings) at what will be the tip 14 of the cantilever 16, which will be used for biofunctionalization along with alignment marks (not shown in drawings). The second step is a lithography step which  
5 is performed to mask the region which will comprise the cantilever 16 with chrome layer 178 (not shown in drawings).

The device 10 is suspended by means of a vertical plasma etch ( $\text{NF}_3$ ,  $\text{Cl}_2$ , Ar) which removes the unmasked portions of the membrane defined in layers 150 and 148 resulting in a cantilever 16 as depicted in Fig. 13i and in top plan view as shown in Fig.  
10 13j. A wet etch is then used to remove the chromium mask 178 (not shown in drawings) and the sample is dried with a critical point dryer. The wafer is diced as shown in the top plan view of Fig. 13k.

The microfluidic channels 52 are then fabricated from a silicone elastomer using a patterned photoresist as a mold, which photoresist is then etched away to define the  
15 actual flow channels defined in a molded elastomeric encapsulating body 182 as shown in Fig. 13l. The flow channels 52 are self-sealing with the silicon of device 10 when placed in an 85°C oven for 24 hours. The device 10 or gold pad 180 is then biofunctionalized by conventional means as shown in Fig. 13m, such as by flowing a fluid through flow channels 52 which fluid carries receptor molecules which  
20 preferentially attach to gold pad 180.

#### *Damping Due to Fluff Ball*

A fluffball 60, which is nothing more than a large dissipative molecule, is provided with linking receptor molecules 62. The fluffball 60 and linking molecule 62 are free  
25 floating in the liquid. Linking molecule 62 is adapted to link to a ligand of interest. Tip 14

is also biofunctionalized with receptors adapted to link to the ligand. Capture of the receptors on tip 14 of the ligand of interest with the receptor 62 and fluffball 60 attached will cause the damping coefficient of tip 14 to dramatically increase.

Fig. 14 is a mathematical model of a cantilever 16 of mass M having a fluff ball 60 attached to its tip 14 by a molecule 62 treated as a spring. The fluffball 60 is devised to maximize dissipation and hence noise and is comprised of a star dendrimer. Molecule 62 may be comprised of an alkane or a ligand chain. The equation for the system of Fig. 14 is:

$$M\ddot{x} + \gamma\dot{x} + \kappa x = k_m(x_d - x) + F$$

$$\gamma_d \dot{x}_d = -k_m(x_d - x)$$

$$x_d = \frac{k_m x}{i\omega \gamma_d + k_m}$$

where M is the mass of cantilever 16,  $\gamma$  is the fluidic damping coefficient for cantilever 16,  $\kappa$  the spring constant of the cantilever 16,  $x$  the displacement of the cantilever tip 14,  $k_m$  the effective spring constant of molecule 62 and  $x_d$  the displacement of fluffball 60,  $\gamma_d$  is the fluidic damping coefficient for fluffball 60, and  $F$  is the external force applied to cantilever 16.

The equation of motion for this system can be rewritten to show that there is an effective damping and effective spring constant of the system given by:

$$\bar{\gamma} = \gamma + \frac{\gamma_d}{1 + \left(\frac{\gamma_d}{k_m} \omega\right)^2}$$

$$\bar{\kappa} = \kappa + k_m \frac{\left(\frac{\gamma_d \omega}{k_m}\right)^2}{1 + \left(\frac{\gamma_d \omega}{k_m}\right)^2}$$

The fluffball 60 is chosen so that its damping coefficient is as large as possible in order to maximize dissipation and hence the noise so that

$$\gamma_d = k_m / \omega$$

$$\bar{\gamma} = \gamma + \frac{1}{2} \frac{k_m}{\omega}$$

- 5 The fractional dissipative effect of the biomolecule and fluffball on dissipation and noise is of the order of:

$$\frac{\Delta\gamma}{\gamma} \approx \frac{k_m}{\gamma \omega}$$

- 10 Many alterations and modifications may be made by those having ordinary skill in the art without departing from the spirit and scope of the invention. Therefore, it must be understood that the illustrated embodiment has been set forth only for the purposes of example and that it should not be taken as limiting the invention as defined by the following claims. For example, notwithstanding the fact that the elements of a claim are
- 15 set forth below in a certain combination, it must be expressly understood that the invention includes other combinations of fewer, more or different elements, which are disclosed in above even when not initially claimed in such combinations.

- The words used in this specification to describe the invention and its various embodiments are to be understood not only in the sense of their commonly defined
- 20 meanings, but to include by special definition in this specification structure, material or acts beyond the scope of the commonly defined meanings. Thus if an element can be understood in the context of this specification as including more than one meaning, then its use in a claim must be understood as being generic to all possible meanings supported by the specification and by the word itself.



The definitions of the words or elements of the following claims are, therefore, defined in this specification to include not only the combination of elements which are literally set forth, but all equivalent structure, material or acts for performing substantially the same function in substantially the same way to obtain substantially the same result. In this sense it is therefore contemplated that an equivalent substitution of two or more elements may be made for any one of the elements in the claims below or that a single element may be substituted for two or more elements in a claim. Although elements may be described above as acting in certain combinations and even initially claimed as such, it is to be expressly understood that one or more elements from a claimed combination can in some cases be excised from the combination and that the claimed combination may be directed to a subcombination or variation of a subcombination.

Insubstantial changes from the claimed subject matter as viewed by a person with ordinary skill in the art, now known or later devised, are expressly contemplated as being equivalently within the scope of the claims. Therefore, obvious substitutions now or later known to one with ordinary skill in the art are defined to be within the scope of the defined elements.

The claims are thus to be understood to include what is specifically illustrated and described above, what is conceptionally equivalent, what can be obviously substituted and also what essentially incorporates the essential idea of the invention.

We claim:

- 1 1. A submicron bioNEMS device comprising:  
2 a support; and  
3 a piezoresistive cantilever coupled to the support extending therefrom with a  
4 length  $l$  and having a width  $w$  and a tip, wherein the cantilever has a restriction portion  
5 of reduced width,  $b$ , and a length  $l_1$ , and a biofunctionalized portion at or near the tip.
- 1 2. The bioNEMS device of claim 1 where the restriction portion is comprised of  
2 multiple legs of reduced width,  $b$ , attached to the support.
- 1 3. The bioNEMS device of claim 2 where the multiple legs are two in number and  
2 are separated from each other by a distance of  $w - 2b$ .
- 1 4. The bioNEMS device of claim 1 further comprising a source of bias current  
2 applied to the restriction portion of the cantilever and where the magnitude of the bias  
3 current is limited by a maximal acceptable temperature increase at the biofunctionalized  
4 tip.
- 1 5. The bioNEMS device of claim 4 where the maximal acceptable temperature  
2 increase at the biofunctionalized tip is approximately 1 degree K.

1 6. An improvement in a piezoresistive bioNEMS device immersed in a fluid  
2 comprising at least one oscillating cantilever having a length  $l$  having a magnitude  
3 chosen to minimize background Johnson noise relative to signal strength generated by  
4 the piezoresistive bioNEMS device.

1 7. The improvement of claim 6 wherein the signal strength is based on  
2 thermomechanical noise levels of the piezoresistive bioNEMS device in the fluid.

1 8. The improvement of claim 6 where the piezoresistive cantilever has a width  $w$ ,  
2 and a restriction portion of reduced width,  $b$ , wherein the reduced width,  $b$ , is chosen to  
3 reduce Johnson noise relative to the signal strength generated by the piezoresistive  
4 bioNEMS device.

1 9. The improvement of claim 8 wherein the signal strength is based on  
2 thermomechanical noise levels of the piezoresistive bioNEMS device in the fluid.

1 10. An improvement in a biofunctionalized bioNEMS device immersed in a fluid  
2 comprising a receptor disposed on the bioNEMS device for binding to a ligand of  
3 interest and a catalyst disposed on the bioNEMS device with the receptor to enhance  
4 binding rate coefficients of the receptor with the ligand of interest.

1 11. The improvement of claim 10 where the catalyst lowers the receptor-ligand  
2 binding activation energy.

1 12. The improvement of claim 10 wherein the receptor is designed by forced  
2 evolution to preferentially bind with the ligand of interest.

1 13. A submicron device comprising:  
2 a source of a carrier signal;  
3 a support;  
4 a piezoresistive cantilever coupled to the support and extending therefrom; and  
5 an element disposed on the cantilever and electromagnetically coupled to the  
6 source so that the cantilever is driven by the carrier signal from the source.

1 14. The device of claim 13 where the element comprises a magnetic film disposed  
2 on the cantilever, and where the source generates an electromagnetic signal coupled to  
3 the magnetic film.

1 15. An apparatus comprising:  
2 a plurality of NEMS transducers, each of the NEMS transducers generating an  
3 output signal; and  
4 means for processing the plurality of corresponding output signals from the  
5 plurality of NEMS transducers to obtain a collective output signal.

1 16. The apparatus of claim 15 where the means averages the plurality of output  
2 signals so that the collective output signal is an average.

1 17. The apparatus of claim 15 where the means determines if a predetermined  
2 fraction of the plurality of output signals are above a threshold within a predetermined  
3 time window.

1 18. The apparatus of claim 15 where each of the plurality of NEMS transducers are  
2 biofunctionalized and where the means effectively increases ligand capture rates by  
3 only generating a collective output signal indicative of an increase ligand capture rate  
4 as compared to a single one of the NEMS transducers.

1 19. An apparatus immersed in a fluid comprising a plurality of NEMS transducers  
2 immersed in the fluid forming an array of adjacent transducers, each of the NEMS  
3 transducers generating an output signal, the motion of two adjacent NEMS transducers  
4 being coupled to each through the fluid in which the adjacent NEMS transducers are  
5 immersed.

1 20. The apparatus of claim 19 where a cross correlation of movement of a first and  
2 second NEMS transducer comprising the two adjacent NEMS transducers,  $C_{12} = \langle x_1$   
3  $(0) x_2(t) \rangle$ , is defined by

$$4 \quad \frac{d}{dt} C_{12}(t) = -k_B T X_{12}(t) \quad \text{for } t > 0$$

5 where  $k_B$  is the Boltzman constant,  $T$  is the temperature of the fluid,  $t$  is time, and  $X_{12}$   
6 the "susceptibility" giving the displacement  $x_2(t)$  of the second transducer for a force  $F_1$   
7 acting on the first transducer, such that an ensemble average for the position of the  
8 second transducer is defined by

9 
$$\langle x_2(t) \rangle = \int_{-\infty}^{\infty} X_{12}(t-t') F_1(t') dt'.$$

1 21. An apparatus immersed in a fluid comprising:  
2 a microfluidic flow channel for carrying a flow of the fluid; and  
3 at least one NEMS transducer disposed in the microfluidic flow channel so that a  
4 characteristic of the fluid is sensed by the NEMS transducer.

1 22. The apparatus of claim 21 where the NEMS transducer is biofunctionalized and  
2 the characteristic of the fluid is sensed by the NEMS transducer is the presence or  
3 absence within the fluid of a ligand to which the NEMS transducer has been  
4 biofunctionalized.

1 23. The apparatus of claim 21 further comprising a plurality of the NEMS  
2 transducers, each of which is disposed in common in the flow channel.

1 24. The apparatus of claim 23 further comprising a plurality of flow channels among  
2 which the plurality of NEMS transducers are distributed.

1 25. The apparatus of claim 23 where the plurality of NEMS transducers are surface  
2 fabricated.

1 26. The apparatus of claim 23 where the plurality of NEMS transducers are  
2 membrane fabricated.

1 27. A method of fabricating a bioNEMS device from a membrane comprising:  
2 providing a heterostructure comprising a wafer layer, an etch stop layer on the  
3 wafer layer, a NEMS device layer on the etch stop layer, and a piezoresistive layer on  
4 the NEMS device layer;  
5 etching trenches through the wafer layer to the etch stop layer to define an area  
6 which will become a membrane in which the NEMS device will be defined;  
7 removing the etch stop layer in the bottom of the trenches to the device layer to  
8 form the membrane;  
9 selectively forming conductive contacts on the piezoresistive layer of the  
10 membrane by electron beam lithography;  
11 selectively forming regions which will become biofunctionalized on the  
12 piezoresistive layer of the membrane by electron beam lithography;  
13 selectively forming a NEMS device on the piezoresistive layer of the membrane  
14 by electron beam lithography which include the region which will become  
15 biofunctionalized;  
16 selectively plasma etching the membrane to remove unmasked portions to  
17 define a suspended NEMS device;  
18 selectively molding a flow channel in an elastomeric layer disposed around the  
19 membrane; and  
20 biofunctionalizing selected regions on the NEMS device.

1 28. The method of claim 27 where providing a heterostructure further comprises  
2 polishing the wafer layer to promote adhesion to the elastomeric layer.

1 29. The method of claim 27 where providing a heterostructure further comprises  
2 thinning the wafer layer.

1 30. The method of claim 27 where selectively plasma etching the membrane to  
2 remove unmasked portions to define a suspended NEMS device comprises selectively  
3 vertically plasma etching away unmasked portions of the NEMS device layer.

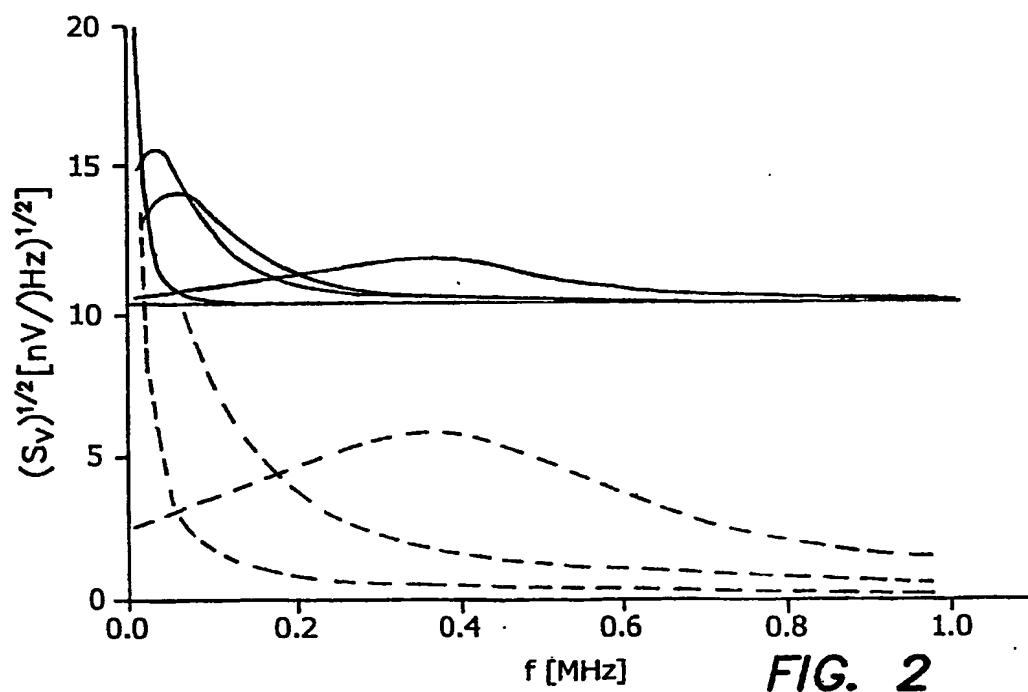
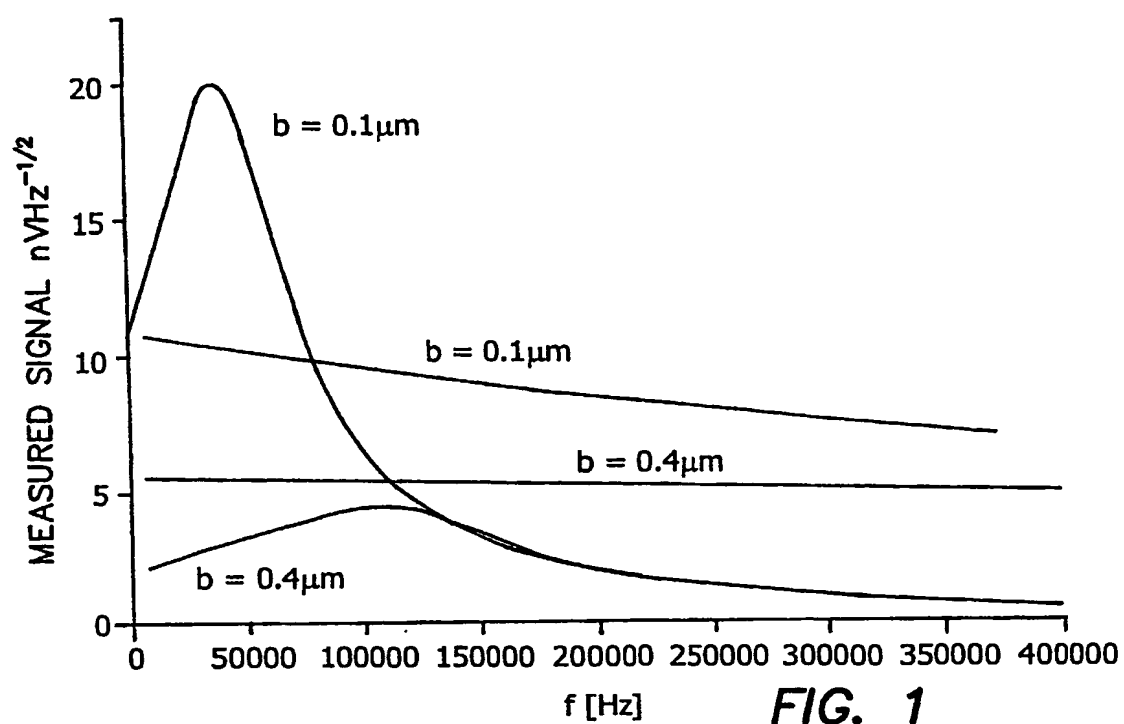
1 31. The method of claim 27 where selectively molding a flow channel in an  
2 elastomeric layer disposed around the membrane comprises selectively disposing a  
3 photoresist layer to define the flow channel, disposing an elastomeric layer on the  
4 selectively disposed photoresist layer, and removing the photoresist layer to define the  
5 flow channel.

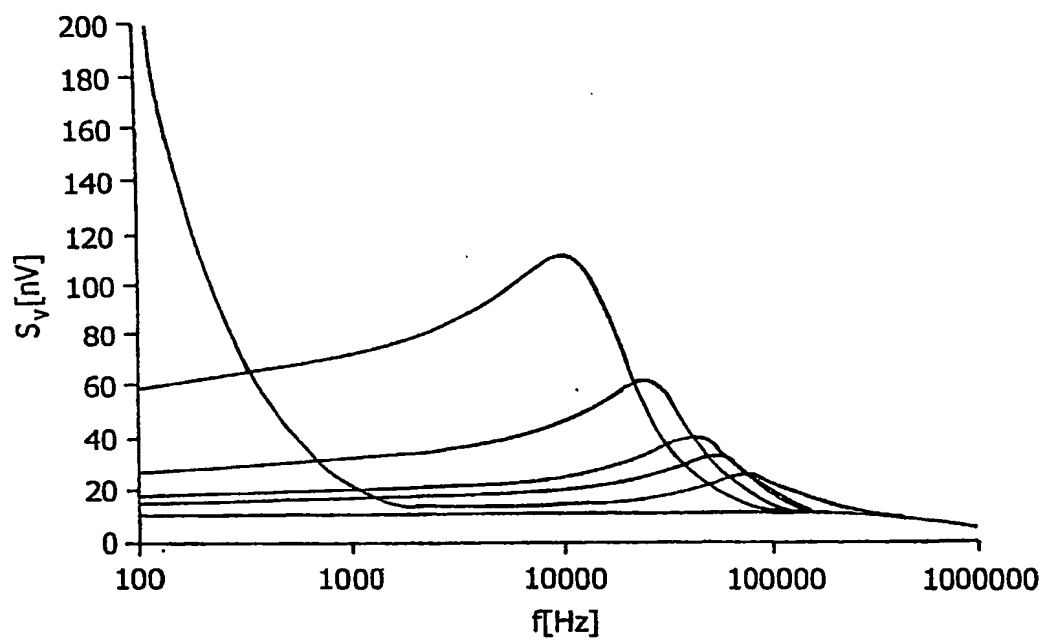
1 32. An NEMS device immersed in a fluid for detecting a ligand comprising:  
2 a resonating member having a tip immersed in the fluid biofunctionalized for the  
3 ligand;  
4 a linking molecule disposed in the fluid; and  
5 a fluffball for providing a damping force disposed in the fluid, wherein capture of  
6 the ligand by the linking molecule, and capture of the fluffball by the linking molecule  
7 causes increased damping of the member when the ligand is attached to the  
8 biofunctionalized tip of the member.

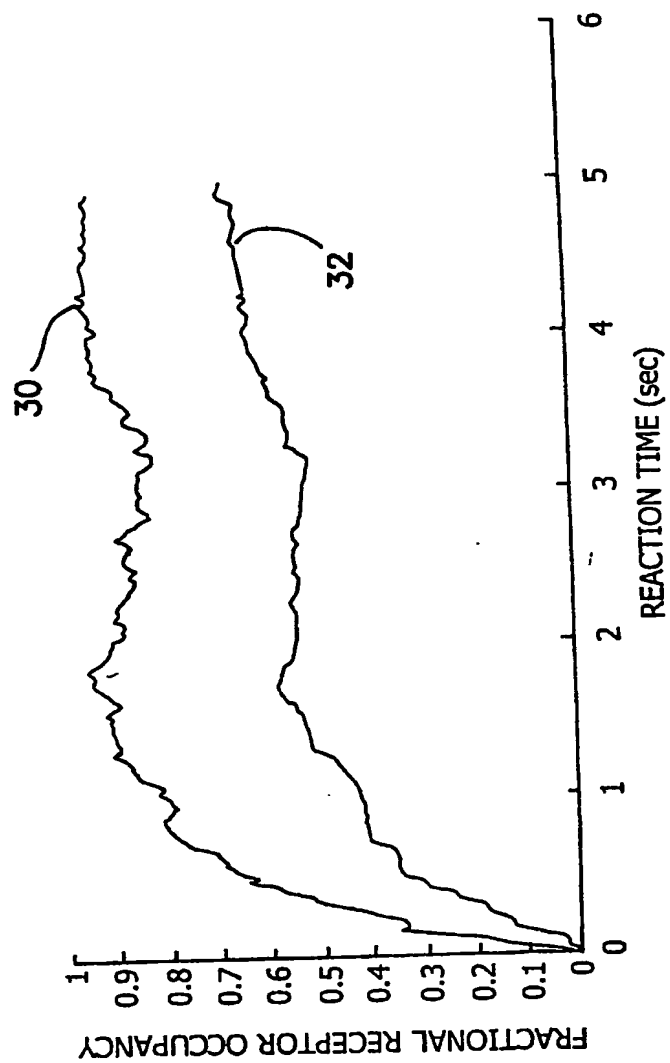
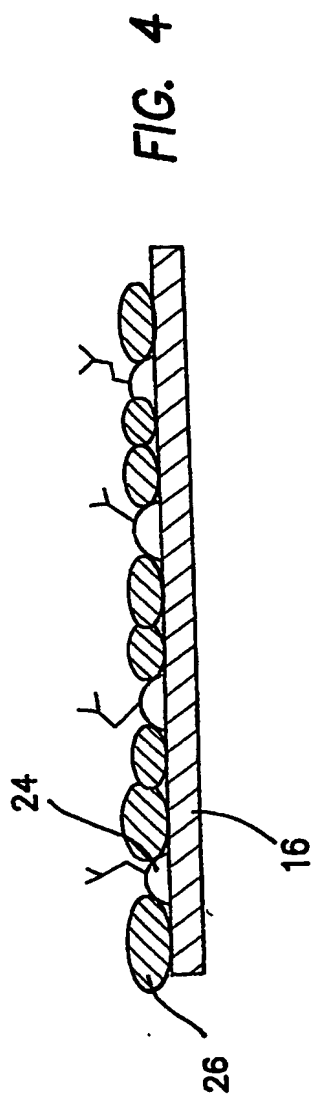
1 33. The NEMS device of claim 32 where the fluffball is comprised of star dendrimer.



- 1 34. A method of detecting a ligand in a fluid by means of a NEMS device comprising:  
2 immersing the tip of a resonating member in the fluid which tip is  
3 biofunctionalized for the ligand;  
4 providing a linking molecule disposed in the fluid;  
5 providing a fluffball in the fluid;  
6 capturing the ligand by the linking molecule;  
7 capturing the fluffball by the linking molecule; and  
8 linking the ligand to the biofunctionalized tip of the resonating member so that  
9 damping of the resonating member is increased.



**FIG. 3**



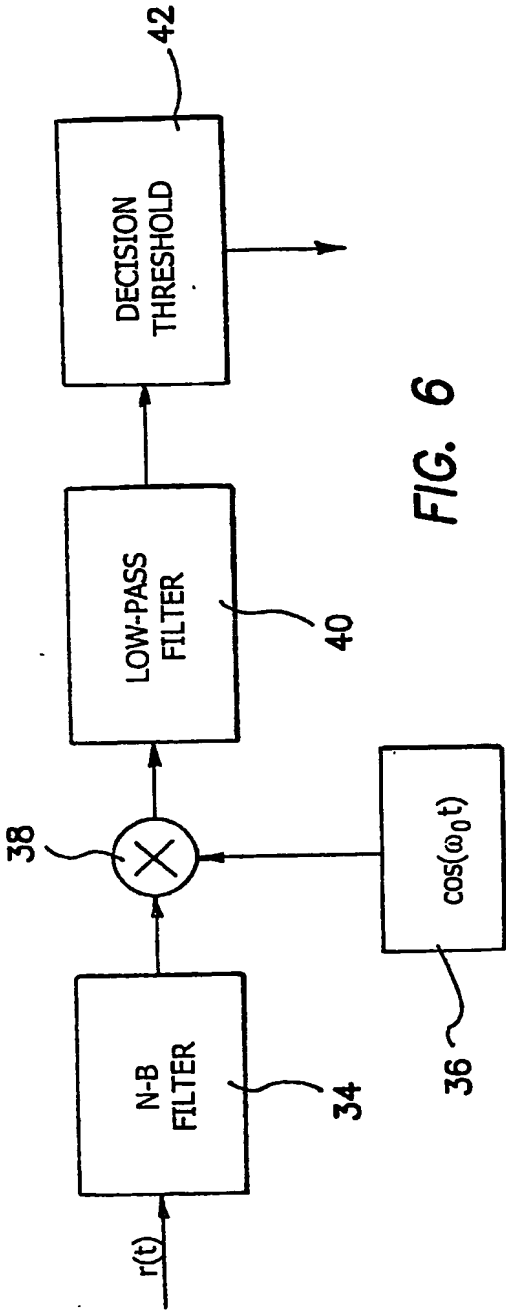


FIG. 6

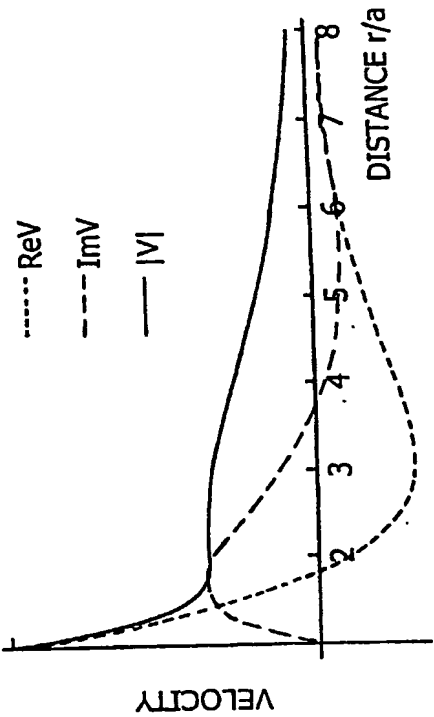
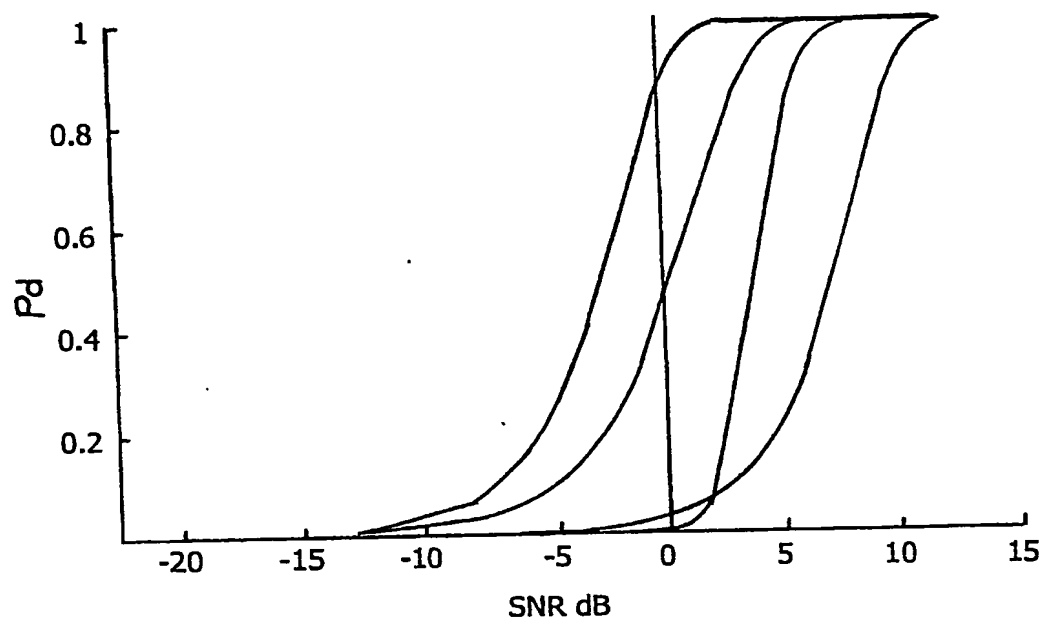
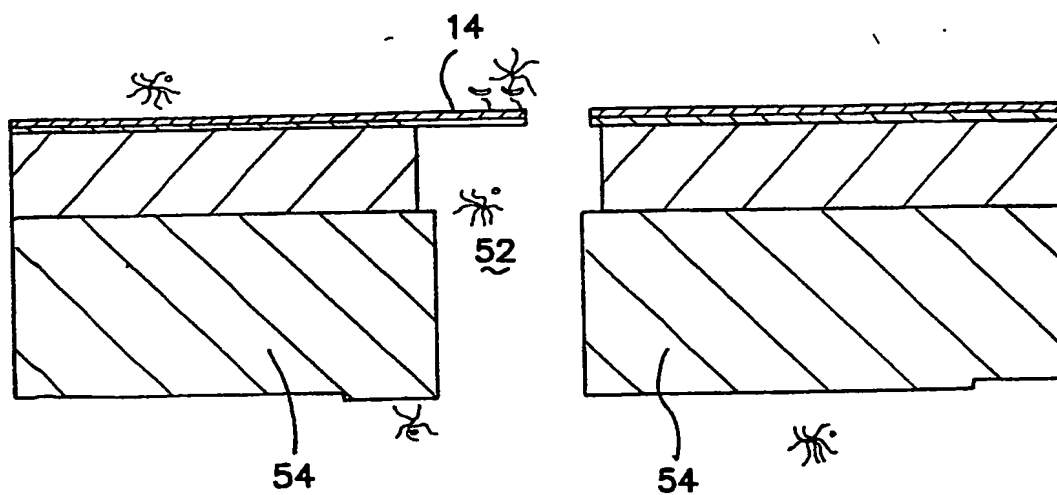


FIG. 9

**FIG. 7****FIG. 10**

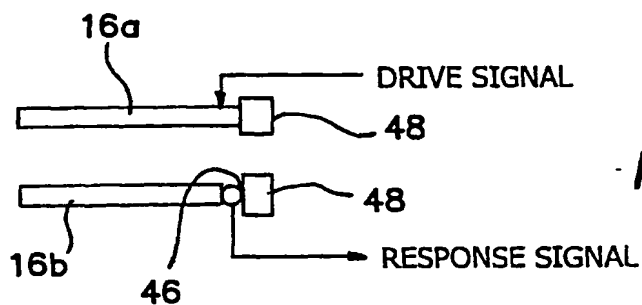


FIG. 8A

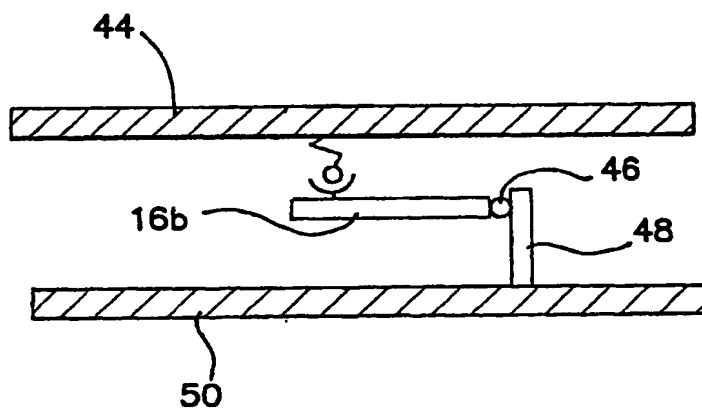


FIG. 8B

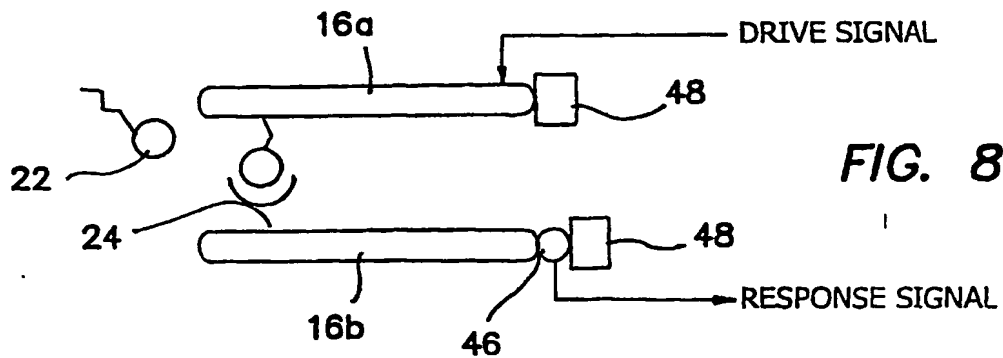


FIG. 8C

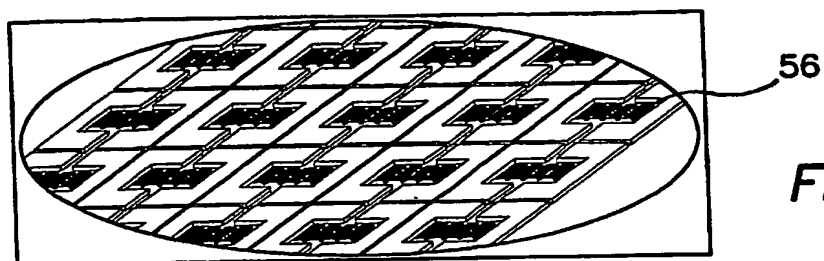


FIG. 11A

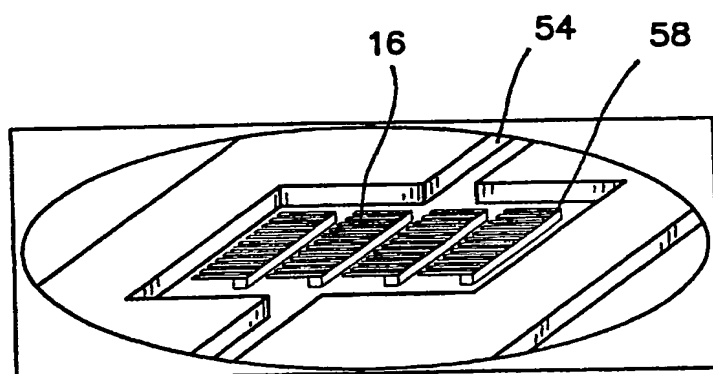


FIG. 11B

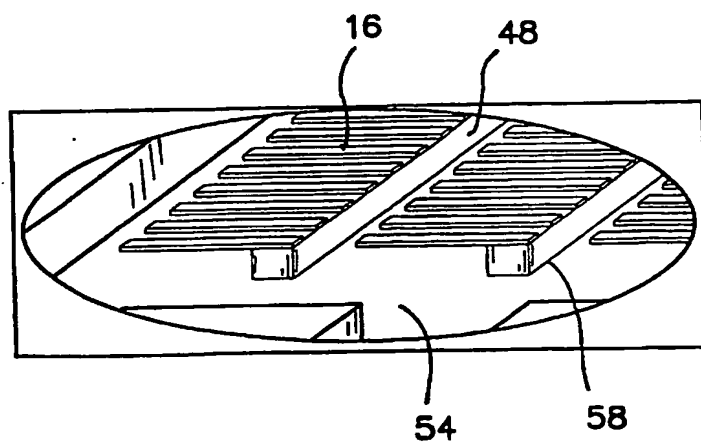
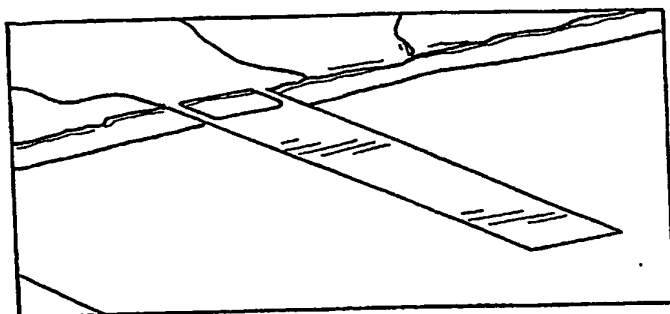


FIG. 11C





10

FIG. 12A

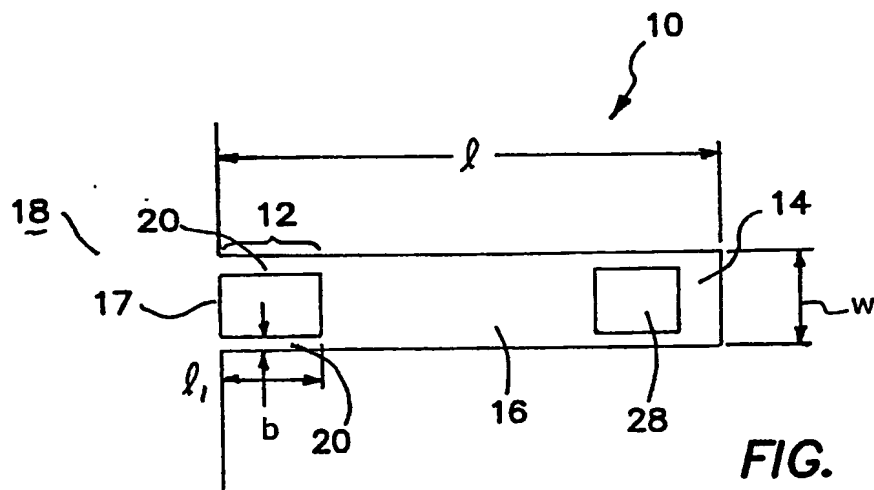


FIG. 12B

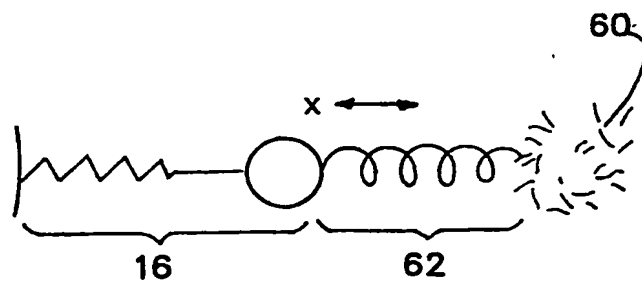


FIG. 14

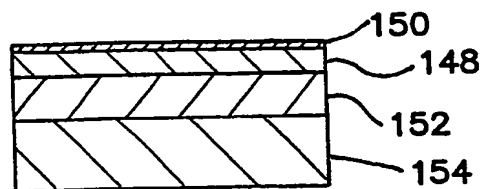


FIG. 13A

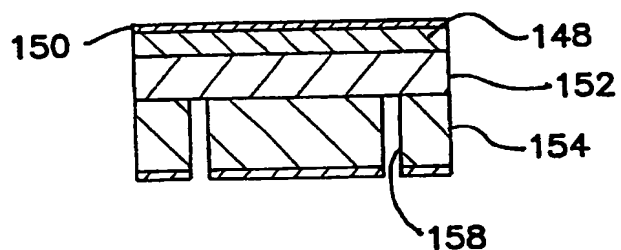


FIG. 13B

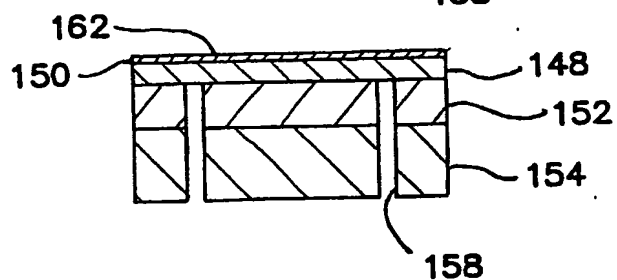


FIG. 13C

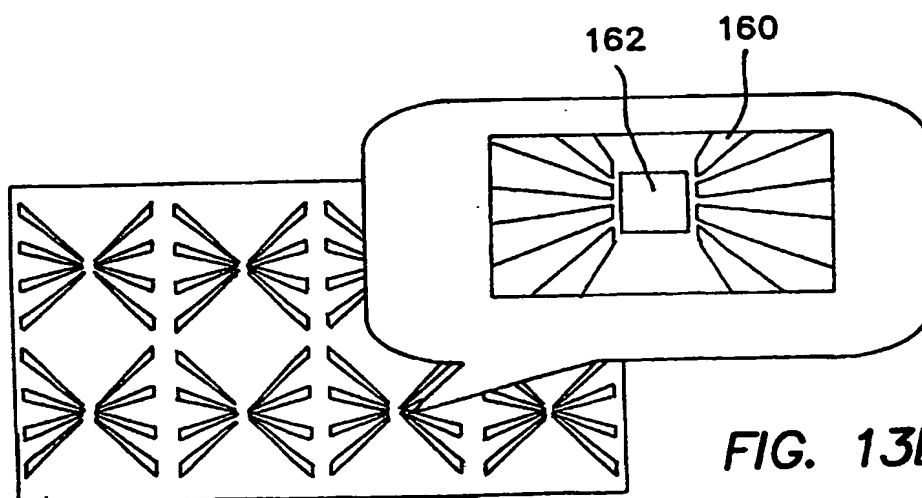


FIG. 13D

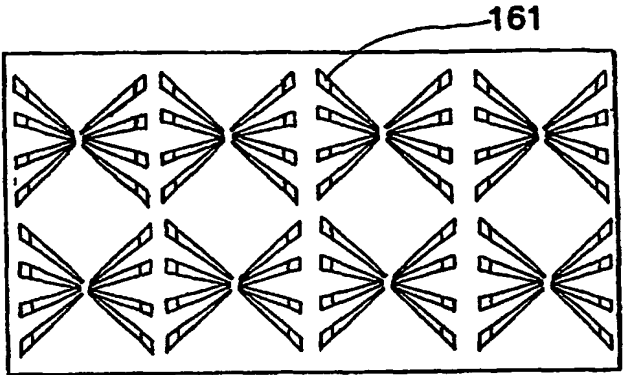


FIG. 13E

FIG. 13F

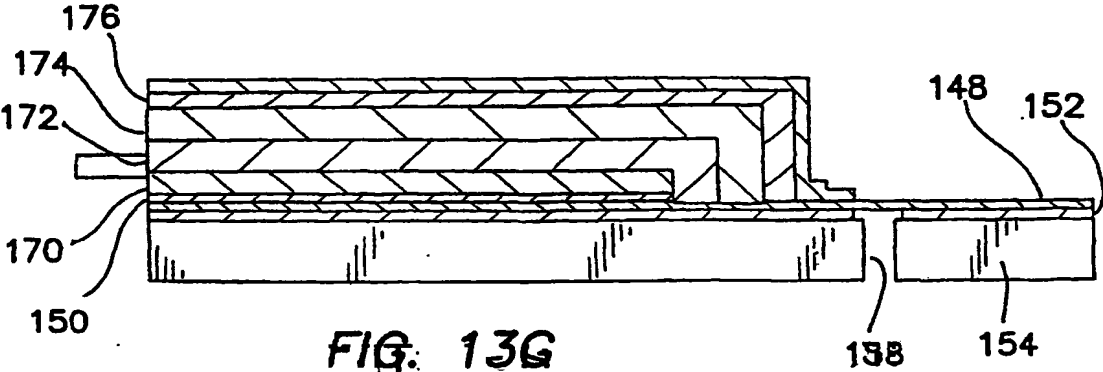
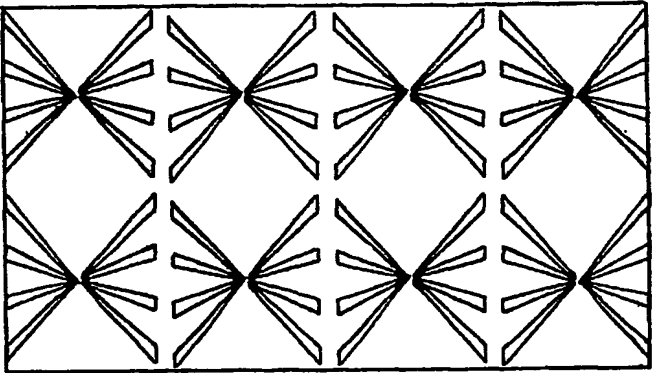


FIG. 13G

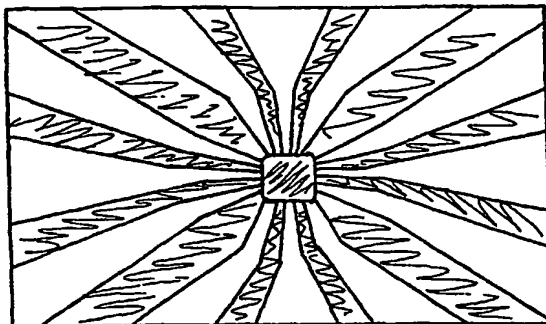


FIG. 13H

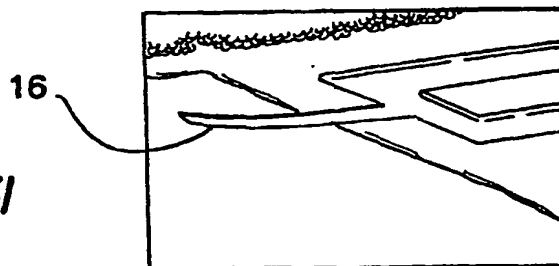


FIG. 13I

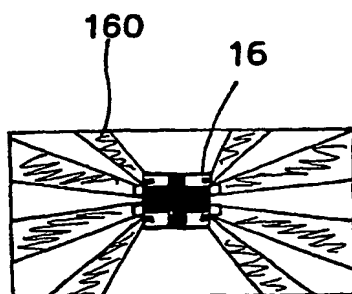
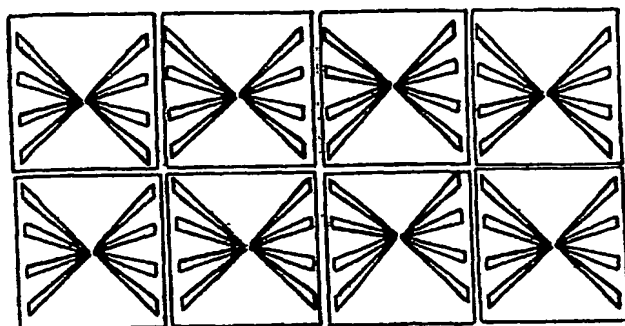
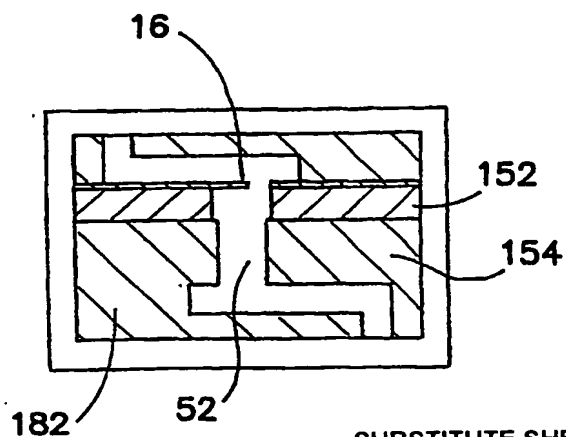
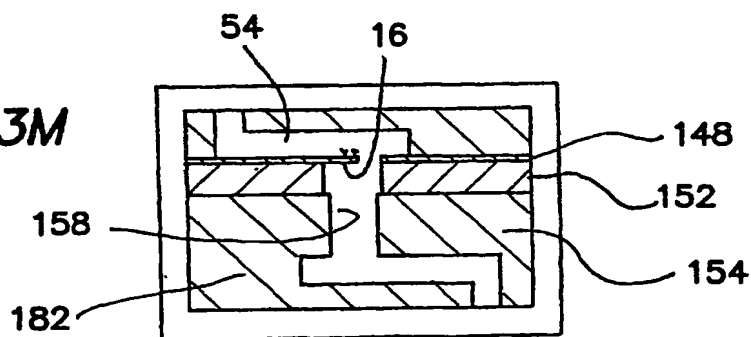


FIG. 13J



**FIG. 13K**

**FIG. 13M**



**FIG. 13L**